

SENATE BILL No. 305

DIGEST OF INTRODUCED BILL

Citations Affected: IC 16-42-22.

Synopsis: Substitution of antiepileptic drug. Prohibits a pharmacist from substituting another brand name or generic antiepileptic drug unless the pharmacist receives consent from specified individuals.

Effective: July 1, 2009.

Miller

January 7, 2009, read first time and referred to Committee on Health and Provider Services.

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First Regular Session 116th General Assembly (2009)

PRINTING CODE. Amendments: Whenever an existing statute (or a section of the Indiana Constitution) is being amended, the text of the existing provision will appear in this style type, additions will appear in **this style type**, and deletions will appear in ~~this style type~~.

Additions: Whenever a new statutory provision is being enacted (or a new constitutional provision adopted), the text of the new provision will appear in **this style type**. Also, the word **NEW** will appear in that style type in the introductory clause of each SECTION that adds a new provision to the Indiana Code or the Indiana Constitution.

Conflict reconciliation: Text in a statute in *this style type* or ~~this style type~~ reconciles conflicts between statutes enacted by the 2008 Regular Session of the General Assembly.

SENATE BILL No. 305

A BILL FOR AN ACT to amend the Indiana Code concerning health.

Be it enacted by the General Assembly of the State of Indiana:

1 SECTION 1. IC 16-42-22-8, AS AMENDED BY P.L.204-2005,
2 SECTION 10, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
3 JULY 1, 2009]: Sec. 8. (a) **Except as provided in subsection (c)**, for
4 substitution to occur for a prescription other than a prescription filled
5 under the Medicaid program (42 U.S.C. 1396 et seq.), the children's
6 health insurance program established under IC 12-17.6-2, or the
7 Medicare program (42 U.S.C. 1395 et seq.):

8 (1) the practitioner must:

9 (A) sign on the line under which the words "May substitute"
10 appear; or

11 (B) for an electronically transmitted prescription,
12 electronically transmit the instruction "May substitute."; and

13 (2) the pharmacist must inform the customer of the substitution.

14 (b) This section does not authorize any substitution other than
15 substitution of a generically equivalent drug product.

16 **(c) A pharmacist may not substitute another brand name or a**
17 **generically equivalent drug product for a prescribed antiepileptic**



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1 **drug unless the pharmacist receives the consent of:**

2 **(1) the customer's prescribing physician; and**

3 **(2) the customer or the customer's legal guardian.**

4 SECTION 2. IC 16-42-22-10, AS AMENDED BY P.L.204-2005,
5 SECTION 12, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
6 JULY 1, 2009]: Sec. 10. (a) If a prescription is filled under the
7 Medicaid program (42 U.S.C. 1396 et seq.), the children's health
8 insurance program established under IC 12-17.6-2, or the Medicare
9 program (42 U.S.C. 1395 et seq.), the pharmacist shall substitute a
10 generically equivalent drug product and inform the customer of the
11 substitution if the substitution would result in a lower price unless:

12 (1) the words "Brand Medically Necessary" are:

13 (A) written in the practitioner's own writing on the form; or

14 (B) electronically transmitted with an electronically
15 transmitted prescription; **or**

16 (2) the practitioner has indicated that the pharmacist may not
17 substitute a generically equivalent drug product by:

18 (A) orally stating that a substitution is not permitted; or

19 (B) for an electronically transmitted prescription, indicating
20 with the electronic prescription that a substitution is not
21 permitted; **or**

22 **(3) the prescribed drug is an antiepileptic drug.**

23 (b) If a practitioner orally states that a generically equivalent drug
24 product may not be substituted, the practitioner must subsequently
25 forward to the pharmacist a written or electronically transmitted
26 prescription with the "Brand Medically Necessary" instruction
27 appropriately indicated in the physician's own handwriting.

28 (c) This section does not authorize any substitution other than
29 substitution of a generically equivalent drug product.

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